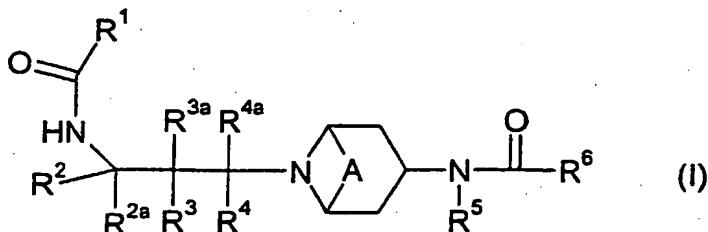


CLAIMS

1. A compound of formula (I):



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wherein:

A is CH_2CH_2 or A is absent;

R^1 is C_{3-7} cycloalkyl (substituted by one or two fluorine atoms and optionally further substituted by C_{1-4} alkyl) or N-linked heterocyclyl (substituted by one or two fluorine atoms and optionally further substituted by C_{1-4} alkyl);

10 R^2 is C_{3-6} alkyl or C_{3-6} cycloalkyl, or phenyl or heteroaryl either of which is optionally substituted by halogen, C_{1-4} alkyl, C_{1-4} alkoxy, $\text{S}(\text{O})_n(\text{C}_{1-4}$ alkyl), nitro, cyano or CF_3 ;

R^{2a} , R^4 and R^{4a} are, independently, hydrogen or C_{1-4} alkyl;

R^3 and R^{3a} are, independently, hydrogen or C_{1-4} alkyl or C_{1-4} alkoxy;

15 R^5 is hydrogen, C_{1-4} alkyl (optionally substituted by halogen, hydroxy, C_{1-4} alkoxy, C_{3-7} cycloalkyl, SH , C_{1-4} alkylthio, cyano or $\text{S}(\text{O})_q(\text{C}_{1-4}$ alkyl)), C_{3-4} alkenyl, C_{3-4} alkynyl or C_{3-7} cycloalkyl;

20 R^6 is phenyl, heteroaryl, phenylNH, heteroarylNH, phenyl(C_{1-2})alkyl, heteroaryl(C_{1-2})alkyl, phenyl(C_{1-2} alkyl)NH or heteroaryl(C_{1-2} alkyl)NH;

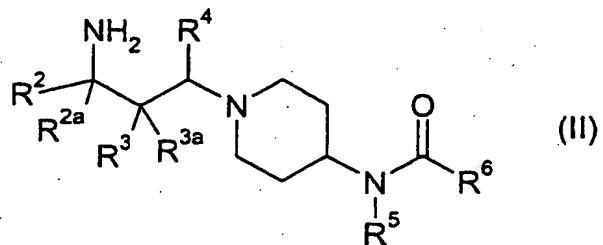
wherein the phenyl and heteroaryl rings of any of the foregoing are, unless stated otherwise, independently optionally substituted by halo, cyano, nitro, hydroxy, C_{1-4} alkyl, C_{1-4} alkoxy, $\text{S}(\text{O})_m\text{C}_{1-4}$ alkyl, $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$, $\text{NHS}(\text{O})_2(\text{C}_{1-4}$ alkyl), NH_2 , $\text{NH}(\text{C}_{1-4}$ alkyl), $\text{N}(\text{C}_{1-4}$ alkyl)₂, $\text{NHC}(\text{O})\text{NH}_2$, $\text{C}(\text{O})\text{NH}_2$, $\text{C}(\text{O})\text{NH}(\text{C}_{1-4}$ alkyl), $\text{NHC}(\text{O})(\text{C}_{1-4}$ alkyl), CO_2H , $\text{CO}_2(\text{C}_{1-4}$ alkyl), $\text{C}(\text{O})(\text{C}_{1-4}$ alkyl), CF_3 , CHF_2 , CH_2F , CH_2CF_3 or OCF_3 ;

25 R^7 and R^8 are, independently, hydrogen or C_{1-4} alkyl, or together with a nitrogen or oxygen atom, may join to form a 5- or 6-membered ring which is optionally substituted with C_{1-4} alkyl, $\text{C}(\text{O})\text{H}$ or $\text{C}(\text{O})(\text{C}_{1-4}$ alkyl);

m , n and q are, independently, 0, 1 or 2;

or a pharmaceutically acceptable salt thereof or a solvate thereof.

2. A compound as claimed in claim 1 wherein R^{2a}, R³, R^{3a} and R⁴ are all hydrogen.
3. A compound as claimed in claim 1 or 2 wherein R^{4a} is hydrogen or methyl.
- 5 4. A compound as claimed in claim 1, 2 or 3 wherein R¹ is C₃₋₇ cycloalkyl (substituted by 1 or 2 fluorine atoms and optionally further substituted by C₁₋₄ alkyl).
- 10 5. A compound as claimed in claim 1, 2, 3 or 4 wherein R¹ is 4,4-di-fluoro-cyclohexyl, 3,3-di-fluoro-cyclopentyl or 3,3-di-fluoro-cyclobutyl.
6. A compound as claimed in claim 1, 2, 3, 4 or 5 wherein R² is phenyl or 6-membered heteroaryl optionally substituted by halogen or CF₃.
- 15 7. A compound as claimed in claim 1, 2, 3, 4, 5 or 6 wherein R⁵ is ethyl.
8. A compound as claimed in claim 1, 2, 3, 4, 5, 6 or 7 wherein R⁶ is phenyl, heteroaryl, phenylNH, heteroarylNH, phenyl(C₁₋₂)alkyl, heteroaryl(C₁₋₂)alkyl, phenyl(C₁₋₂ alkyl)NH or heteroaryl(C₁₋₂ alkyl)NH (for example phenyl or phenylCH₂); wherein the phenyl and heteroaryl rings of R⁶ are substituted by S(O)₂C₁₋₄ alkyl, and optionally further substituted by one or more of halo, cyano, nitro, hydroxy, C₁₋₄ alkyl, C₁₋₄ alkoxy, S(O)_mC₁₋₄ alkyl, S(O)₂NR⁷R⁸, NHS(O)₂(C₁₋₄ alkyl), NH₂, NH(C₁₋₄ alkyl), N(C₁₋₄ alkyl)₂, NHC(O)NH₂, C(O)NH₂, C(O)NH(C₁₋₄ alkyl), NHC(O)(C₁₋₄ alkyl), CO₂H, CO₂(C₁₋₄ alkyl), C(O)(C₁₋₄ alkyl), CF₃, CHF₂, CH₂F, CH₂CF₃ or OCF₃; wherein m, R⁷ and R⁸ are as defined in claim 1.
- 25 9. A process for the preparation of a compound of formula (I) as claimed in claim 1, wherein A is absent, comprising treating a compound of formula (II):



with:

an acid chloride of formula $R^1C(O)Cl$, in the presence of a base and in a suitable solvent; or,

an acid of formula R^1CO_2H , in the presence of a suitable coupling agent, a suitable base and in a suitable solvent.

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10. A pharmaceutical composition which comprises a compound of the formula (I), or a pharmaceutically acceptable salt thereof or solvate thereof as claimed in claim 1, and a pharmaceutically acceptable adjuvant, diluent or carrier.

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11. A compound of the formula (I), or a pharmaceutically acceptable salt thereof or solvate thereof as claimed in claim 1, for use in therapy.

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12. A compound of formula (I), or a pharmaceutically acceptable salt thereof or solvate thereof as claimed in claim 1, in the manufacture of a medicament for use in therapy.

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13. A method of treating a chemokine mediated disease state in a warm blooded animal suffering from, or at risk of, said disease, which comprises administering to an animal in need of such treatment a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt thereof or solvate thereof as claimed in claim 1.

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